## 8.0 BIO-DETEK Electrode Adaptor 510(k) Summary

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| **Company:** | BIO-DETEK<br>525 Narragansett Park Drive<br>Pawtucket, RI 02861-4323 |
| **Contact:** | Robert Morse<br>QA/QC Manager |
| **Date Prepared:** | January 12, 2004 |
| **Name of Device:** | BIO-DETEK Electrode Adaptor |
| **Predicate Device:** | Heartstream Electrode Adaptor K#984286 |

**Device Description and Intended Use:**

The BIO-DETEK Electrode Adaptor is indicated for use with ZOLL CPR-D, Stat*padz, and Stat*padz II Electrodes for connection with Medtronic Physio-Control Defibrillator Models: LIFEPAK 9, LIFEPAK 10C, LIFEPAK 11, LIFEPAK 12, and LIFEPAK 20, all fitted with QUIK-COMBO™ Therapy Cable.

The BIO-DETEK Electrode Adaptor is made of rigid thermoplastic and conductive material. The adaptor will be provided as a reusable stand-alone accessory to be used in conjunction with ZOLL CPR-D, Stat*padz, and Stat*padz II Electrodes.

**Technological Characteristics**

The BIO-DETEK Electrode Adaptor is designed to comply with the applicable portions of the following standards:

- IEC 60601-2-4 Medical Electrical Equipment
- ANSI/AAMI/ISO DF39-1993
- ANSI/AAMI EC53:1995
- 21 CFR Par 898 Performance Standards for Electrode Lead Wires and Patient Cables

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

APR - 9 2004

Bio-Detek, Inc.
c/o Mr. Robert Morse
QA/QC Manager
525 Narragansett Park Drive
Pawtucket, RI 02861-4323

Re: K040056
Trade Name: BIO-DETEK Electrode Adaptor
Regulation Number: 21 CFR 870.2360
Regulation Name: Electrocardiograph Electrode
Regulatory Class: III (three)
Product Code: MLN
Dated: January 12, 2004
Received: January 13, 2004

Dear Mr. Morse:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.
This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4648. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/dsma/dsmamain.html

Sincerely yours,

Donna R. Lachman

Bram D. Zuckerman, M.D.
Director
Division of Cardiovascular Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

## Indications for Use

510(k) Number: K040056

Device Name: BIO-DETEK Electrode Adapter

Intended Use:

The BIO-DETEK Electrode Adapter is indicated for use with ZOLL CPR-D, stat•padz™ and stat•padz II™ Electrodes for connection with Medtronic Physio-Control Defibrillator Models: LIFEPAK 9, LIFEPAK 10C, LIFEPAK 11, LIFEPAK 12, and LIFEPAK 20, all fitted with QUIK-COMBO Therapy Cable..

Prescription Use ✓ AND/OR Over-The-Counter Use _____
(Part 21 CFR 801 Subpart D)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Diana R. Lechner
(Division Sign-Off)
Division of Cardiovascular Devices

510(k) Number K040056